## CLAIMS

- A MHC Class I-restricted epitope peptide derived from survivin, said epitope having at
   least one of the following characteristics:
- (i) capable of binding to the Class I HLA molecule to which it is restricted at an affinity as measured by the amount of the peptide that is capable of half maximal recovery of the Class I HLA molecule ( $C_{50}$  value) which is at the most 50  $\mu$ M as determined by the assembly binding assay as described herein,
  - (ii) capable of eliciting INF- $\gamma$ -producing cells in a PBL population of a cancer patient at a frequency of at least 1 per  $10^4$  PBLs as determined by an ELISPOT assay, and/or
- 15 (iii) capable of *in situ* detection in a tumor tissue of CTLs that are reactive with the epitope peptide.
  - 2. A peptide according to claim 1 having a  $C_{50}$  value, which is at the most 30  $\mu M$ .
- 20 3. A peptide according to claim 2 having a  $C_{50}$  value, which is at the most 20  $\mu M$ .
  - 4. A peptide according to claim 1, which is restricted by a MHC Class I HLA-A molecule.
- 5. A peptide according to claim 4, which is restricted by a MHC Class I HLA species selected from the group consisting of HLA-A1, HLA-A2, HLA-A3, HLA-A11 and HLA-A24.
  - 6. A peptide according to claim 5, which is restricted by HLA-A2.
- 7. A peptide according to claim 6, which is selected from the group consisting of
  30 FLKLDRERA (SEQ ID NO:1), TLPPAWQPFL (SEQ ID NO:2), ELTLGEFLKL (SEQ ID NO:3),
  LLLGEFLKL (SEQ ID NO:4) and LMLGEFLKL (SEQ ID NO:5).
  - 8. A peptide according to claim 1, which is restricted by a MHC Class I HLA-B molecule.
- 9. A peptide according to claim 8, which is restricted by a MHC Class I HLA-B species selected from the group consisting of HLA-B7, HLA -B35, HLA -B44, HLA-B8, HLA-B15, HLA-B27 and HLA-B51.
  - 10. A peptide according to claim 9, which is restricted by HLA-B35.

11. A peptide according to claim 10, which is selected from the group consisting of CPTENEPDL (SEQ ID NO:6), EPDLAQCFF (SEQ ID NO:7), CPTENEPDY (SEQ ID NO:8) and EPDLAQCFY (SEQ ID NO:9).

- 12. A peptide according to claim 1 comprising at the most 20 amino acid residues.
- 13. A peptide according to claim 12 that comprises at the most 10 amino acid residues.
- 5 14. A peptide according to claim 1, which is a nonapeptide or a decapeptide.
  - 15. A peptide according to claim 1, which is a native sequence of survivin of a mammal species.
- 10 16. A peptide according to claim 15 that is derived from human survivin.
  - 17. A peptide according to claim 1, which is derived from a native sequence of survivin by substituting, deleting or adding at least one amino acid residue.
- 15 18. A peptide according to any of the preceding claims, which is phosphorylated.
  - 19. A peptide according to claim 18, which comprises Thr34 of the native survivin disclosed in US 6.245.523.
- 20 20. A peptide according to claim 1 comprising, for each specific HLA allele, any of the amino acid residues as indicated in the following table:

HLA al-	Position	Position	Position	Position	Position	Position	C-termi-
lele	1	2	3	5	6	7	nal
HLA-A1		T,S	D,E			L	Υ
HLA-A2		L, M	;		\ <b>V</b>		L,V
HLA-A3		L,V,M	F,Y				K, Y, F
HLA-A11		V,I,F,Y	M,L,F,Y,I				K, R
HLA-A23		I,Y					W,I
HLA-A24		Υ		I,V	F		I,L,F
HLA-A25		M,A,T	I				w
HLA-A26	E,D	V,T,I,L,F			I,L,V		Y,F
HLA-A28	E,D	V,A,L					A,R
HLA-A29		E					Y,L
HLA-A30		Y,L,F,V					Y
HLA-A31			L,M,F,Y				R
HLA-A32		I,L					W
HLA-A33		Y,I,L,V					R
HLA-A34		V,L					R
HLA-A66	E,D	T,V					R,K
HLA-A68	E,D	T,V					R,K
HLA-A69		V,T,A					V,L
HLA-A74		Т					V,L
HLA-B5		A,P	F,Y				I,L

HLA-B7	ı	Р	ı	1		1	L,F
HLA-B8		r	ĸ	K,R			L
HLA-B14		R,K	``	17,17			_ L,V
HLA-B15		Q,L,K,P,					F,Y,W
(B62)		H,V,I,M,					.,.,
(002)		S,T					1
HLA-B17		3,1					L,V
HLA-B27		R					Y, K,F,L
HLA-B35		P					I, L, M, Y
HLA-B37		D,E					I,L,M
HLA-B38		H	D,E				F,L
HLA-B39		R,H	·				L,F
HLA-B40		E	F,I,V				L,V,A,W,
(B60,61)							M,T,R
HLA-B42		L,P					Y,L
HLA-B44		E					F,Y,W
HLA-B46		M,I,L,V					Y,F
HLA-B48		Q,K					L
HLA-B51		A,P,G					F,Y,I,V
HLA-B52		Q	F,Y				I,V
HLA-B53		P					W,F,L
HLA-B54		Р					
HLA-B55		P					A,V
HLA-B56		Р					A,V
HLA-B57		A,T,S					F,W,Y
HLA-B58		A,T,S					F,W,Y
HLA-B67		Р					L
HLA-B73		R					Р
HLA-		A,L	ļ				L
Cw1							
HLA-		A,L					F,Y
Cw2							
HLA-		A,L					L,M
Cw3							
HLA-		Y,P,F					L,M,F,Y
Cw4							LTVV
HLA-							L,I,V,Y
Cw6 HLA-		Y					L,Y,F
i .		1					<b>-</b> ,
Cw6 HLA-		Y					L,I,
Cw8		1					<b>-</b> , -,
HLA-		A,L					L,V
Cw16		/3,5					-, -
CMIO		İ	1	l	l	i	

- 21. A peptide according to claims 1 that is capable of eliciting INF- $\gamma$ -producing cells in a PBL population of a cancer patient at a frequency of at least 10 per 10<sup>4</sup> PBLs.
- 5 22. A peptide according to claim 1, which is capable of eliciting INF- $\gamma$ -producing cells in a PBL population of a patient having a cancer disease where survivin is expressed.
- 23. A peptide according to claim 22 where the cancer disease is selected from the group consisting of a haematopoietic malignancy including chronic lymphatic leukemia and
  10 chronic myeloid leukemia, melanoma, breast cancer, cervix cancer, ovary cancer, lung cancer, colon cancer, pancreas cancer and prostate cancer.
- 24. A peptide according to claim 1, which is capable of eliciting INF-γ -producing cells in a PBL population of a patient having a cancer disease, said INF-γ -producing cells having cytotoxic effect against survivin expressing cells of a cancer cell line, including a cell line selected from the group consisting of the breast cancer cell line MCF-7 and the melanoma cell line FM3.
  - 25. A pharmaceutical composition comprising the peptide according to claim 1.
  - 26. A composition according to claim 25 that comprises a peptide according to claim 4 in combination with a peptide according to claim 8.
- 27. A composition according to claim 26 comprising a peptide according to claim 6 in com-25 bination with a peptide according to claim 10.
  - 28. A composition according to claim 25, which is a vaccine capable of eliciting an immune response against a cancer disease.
- 30 29. A composition according to claim 25, further comprising an immunogenic protein or peptide fragment selected from a protein or peptide fragment not belonging to or derived from the survivin protein family.
- 30. A composition according to claim 29, where the protein or peptide fragment not35 belonging to or derived from the survivin protein family is a protein, or peptide fragment hereof, involved in regulation of cell apoptosis.
- 31. A composition according to claim 29 where the immunogenic protein or peptide fragment selected from a protein or peptide fragment hereof not belonging to or derived from the survivin protein family is Bcl-2 or a peptide fragment hereof.
  - 32. A composition according to claim 25, which is a multiepitope vaccine.

- 33. A composition according to claim 28 where the vaccine is capable of eliciting an immune response against a cancer disease where survivin is expressed.
- 34. A composition according to claim 33 where the cancer disease is selected from the group consisting of a haematopoietic malignancy including chronic lymphatic leukemia and chronic myeloid leukemia, melanoma, breast cancer, cervix cancer, ovary cancer, lung cancer, colon cancer, pancreas cancer and prostate cancer.
- 35. A composition according to claim 33 or 34 where the vaccine elicits the production in the vaccinated subject of effector T-cells having a cytotoxic effect against the cancer cells.
  - 36. A composition for ex vivo or in situ diagnosis of the presence in a cancer patient of survivin reactive T-cells among PBLs or in tumor tissue, the composition comprising a peptide according to claim 1.
  - 37. A diagnostic kit for ex vivo or in situ diagnosis of the presence in a cancer patient of survivin reactive T-cells among PBLs or in tumour tissue comprising a peptide according to claim 1.
- 20 38. A complex of a peptide according to claims 1 and a Class I HLA molecule or a fragment of such molecule.
  - 39. A complex according to claim 38 which is monomeric.
- 25 40. A complex according to claim 38 which is multimeric.

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- 41. A method of detecting in a cancer patient the presence of survivin reactive T-cells, the method comprising contacting a tumour tissue or a blood sample with a complex according to claim 38 and detecting binding of the complex to the tissue or the blood cells.
- 42. A molecule that is capable of binding specifically to a peptide according to claims 1.
- 43. A molecule according to claim 36 which is an antibody or a fragment hereof.
- 35 44. A molecule that is capable of blocking the binding of a molecule according to claim 42 or 43.
- 45. A method of treating a cancer disease, the method comprising administering to a patient suffering from the disease an effective amount of the composition according to claim 40 25, a molecule according to claim 42 or a molecule according to claim 44.
  - 46. A method according to claim 45 wherein the disease to be treated is a cancer disease where survivin is expressed.

47. A method according to claim 46 wherein the cancer disease is selected from the group consisting of a haematopoietic malignancy including chronic lymphatic leukemia and chronic myeloid leukemia, melanoma, breast cancer, cervix cancer, ovary cancer, lung cancer, colon cancer, pancreas cancer and prostate cancer.

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- 48. A method according to claim 45, which is combined with a further treatment.
- 49. A method according to claim 48 wherein the further treatment is radiotherapy or chemotherapy.